

10/796, 313

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NEWS 2 "Ask CAS" for self-help around the clock  
NEWS 3 Jul 12 BEILSTEIN enhanced with new display and select options, resulting in a closer connection to BABS  
NEWS 4 AUG 02 IFIPAT/IFIUDB/IFICDB reloaded with new search and display fields  
NEWS 5 AUG 02 CPlus and CA patent records enhanced with European and Japan Patent Office Classifications  
NEWS 6 AUG 02 The Analysis Edition of STN Express with Discover! (Version 7.01 for Windows) now available  
NEWS 7 AUG 27 BIOCOMMERCE: Changes and enhancements to content coverage  
NEWS 8 AUG 27 BIOTECHABS/BIOTECHDS: Two new display fields added for legal status data from INPADOC  
NEWS 9 SEP 01 INPADOC: New family current-awareness alert (SDI) available  
NEWS 10 SEP 01 New pricing for the Save Answers for SciFinder Wizard within STN Express with Discover!  
NEWS 11 SEP 01 New display format, HITSTR, available in WPIDS/WPINDEX/WPIX  
NEWS 12 SEP 27 STANDARDS will no longer be available on STN  
NEWS 13 SEP 27 SWETSCAN will no longer be available on STN

NEWS EXPRESS JULY 30 CURRENT WINDOWS VERSION IS V7.01, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 11 AUGUST 2004

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=> file reg  
COST IN U.S. DOLLARS  
  
FULL ESTIMATED COST

10/796,313

FILE 'REGISTRY' ENTERED AT 18:15:50 ON 20 OCT 2004  
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STRUCTURE FILE UPDATES: 19 OCT 2004 HIGHEST RN 765878-56-6  
DICTIONARY FILE UPDATES: 19 OCT 2004 HIGHEST RN 765878-56-6

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when  
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Crossover limits have been increased. See HELP CROSSOVER for details.

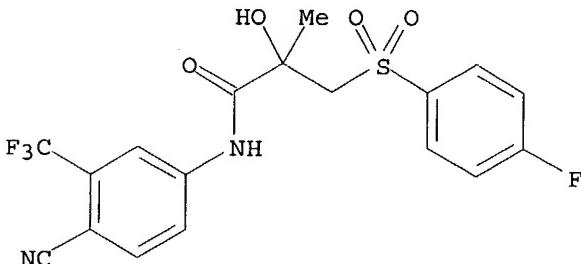
Experimental and calculated property data are now available. For more  
information enter HELP PROP at an arrow prompt in the file or refer  
to the file summary sheet on the web at:  
<http://www.cas.org/ONLINE/DBSS/registryss.html>

```
=> e bicalutamide/cn
E1      1 BICALON/CN
E2      1 BICALPHOS/CN
E3      1 --> BICALUTAMIDE/CN
E4      1 BICARBAMALDEHYDE/CN
E5      1 BICARBAMAMIC ACID/CN
E6      1 BICARBAMAMIDE/CN
E7      1 BICARBAMAMIDE, 2-(4-BIPHENYLYL)-N-METHYL-3-PHENYL-/CN
E8      1 BICARBAMAMIDINE/CN
E9      1 BICARBAMIC ACID/CN
E10     1 BICARBAMIC ACID (4-AMINO-3,5-XYLYL)-, DIETHYL ESTER/CN
E11     1 BICARBAMIC ACID, ((1-CARBOXY-2-HYDROXYETHYL)CARBAMOYL)METHYL
L) -, 1,4-DIBENZYL METHYL ESTER/CN
E12     1 BICARBAMIC ACID, ((CARBOXYMETHYL)CARBAMOYL)METHYL -, DIBENZ
YL METHYL ESTER/CN
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```
=> s e3
L1      1 BICALUTAMIDE/CN
```

```
=> d scan
```

```
L1      1 ANSWERS  REGISTRY  COPYRIGHT 2004 ACS on STN
IN      Propanamide, N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-
fluorophenyl)sulfonyl]-2-hydroxy-2-methyl- (9CI)
MF      C18 H14 F4 N2 O4 S
```



10/796,313

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

ALL ANSWERS HAVE BEEN SCANNED

|                      | SINCE FILE<br>ENTRY | TOTAL<br>SESSION |
|----------------------|---------------------|------------------|
| => file caplus       |                     |                  |
| COST IN U.S. DOLLARS |                     |                  |
| FULL ESTIMATED COST  | 5.27                | 5.48             |

FILE 'CAPLUS' ENTERED AT 18:16:58 ON 20 OCT 2004  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
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FILE COVERS 1907 - 20 Oct 2004 VOL 141 ISS 17  
FILE LAST UPDATED: 19 Oct 2004 (20041019/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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L2      402 L1

=> s l2 and micro?
      2105390 MICRO?
L3      48 L2 AND MICRO?

=> s l3 and particle
      621330 PARTICLE
      701149 PARTICLES
      1052058 PARTICLE
          (PARTICLE OR PARTICLES)
L4      5 L3 AND PARTICLE

=> s l3 and diameter
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          (DIAMETER OR DIAMETERS)
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=> s l2 and micronized
      2954 MICRONIZED
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10/796,313

L6 2 L2 AND MICRONIZED

=> dup rem 16 14

PROCESSING COMPLETED FOR L6

PROCESSING COMPLETED FOR L4

L7 5 DUP REM L6 L4 (2 DUPLICATES REMOVED)

=> d 17 ibib hitstr abs 1-5

L7 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 1

ACCESSION NUMBER: 2004:182593 CAPLUS

DOCUMENT NUMBER: 140:235504

TITLE: Preparation and crystallization of bicalutamide

INVENTOR(S): Dolitzky, Ben-Zion; Reany, Ofer; Shammai, Jenny

PATENT ASSIGNEE(S): Israel

SOURCE: U.S. Pat. Appl. Publ., 12 pp., Cont.-in-part of U.S. Ser. No. 170,721.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

| PATENT NO.             | KIND | DATE     | APPLICATION NO. | DATE        |
|------------------------|------|----------|-----------------|-------------|
| US 2004044249          | A1   | 20040304 | US 2003-606403  | 20030625    |
| US 2003045741          | A1   | 20030306 | US 2002-170721  | 20020613    |
| US 6737550             | B2   | 20040518 |                 |             |
| US 2004059147          | A1   | 20040325 | US 2003-668982  | 20030922    |
| US 6797843             | B2   | 20040928 |                 |             |
| US 2004167349          | A1   | 20040826 | US 2004-791468  | 20040301    |
| US 2004176633          | A1   | 20040909 | US 2004-796313  | 20040308    |
| US 2004176638          | A1   | 20040909 | US 2004-796822  | 20040308    |
| PRIORITY APPLN. INFO.: |      |          | US 2001-298009P | P 20010613  |
|                        |      |          | US 2002-371069P | P 20020409  |
|                        |      |          | US 2002-170721  | A2 20020613 |

OTHER SOURCE(S): CASREACT 140:235504

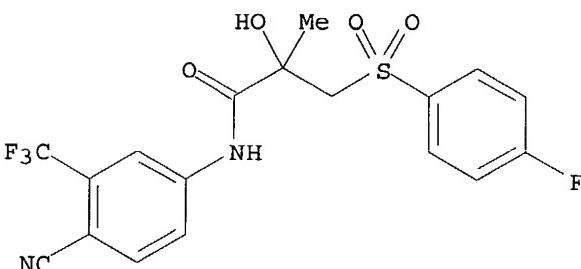
IT 90357-06-5P, Bicalutamide

RL: PEP (Physical, engineering or chemical process); PUR (Purification or recovery); PYP (Physical process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)

(preparation, micronization and crystallization of bicalutamide)

RN 90357-06-5 CAPLUS

CN Propanamide, N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-fluorophenyl)sulfonyl]-2-hydroxy-2-methyl- (9CI) (CA INDEX NAME)



AB Racemic N-[4-cyano-3-trifluoromethylphenyl]-3-[4-fluorophenylsulfonyl]-2-hydroxy-2-methylpropionamide (bicalutamide) was prepared starting from Et pyruvate and Me methacrylate. Thus, 5-amino-2-cyanobenzotrifluoride was treated with DABCO and reacted with deprotonated Et 2-(4-

fluorophenylsulfonyl)-2-hydroxy-2-methylpropionate (prepared from Et pyruvate) to give 40% bicalutamide. Micronized particles of bicalutamide can be obtained as pharmaceutical compns. that are useful for its anti-androgen activity (no data). Bicalutamide intermediates were also prepared, including Et 2-(4-fluorophenylsulfonyl)-2-hydroxy-2-methylpropionate, Me 2,3-epoxy-2-methylpropionate and 2-hydroxy-2-methyl-3-(4-fluorophenylthio)propionic acid. The present invention further discloses the isolation and purification of bicalutamide by various crystallization methods.

L7 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2003:472643 CAPLUS  
 DOCUMENT NUMBER: 139:30801  
 TITLE: Assays and implements for determining and modulating heat shock protein 90 (HSP90) binding activity, and therapeutic use  
 INVENTOR(S): Kamal, Adeela; Burrows, Francis J.; Zhang, Lin; Boehm, Marcus F.  
 PATENT ASSIGNEE(S): Conforma Therapeutics Corporation, USA  
 SOURCE: PCT Int. Appl., 61 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 2  
 PATENT INFORMATION:

| PATENT NO.  | KIND   | DATE     | APPLICATION NO. | DATE     |
|---|--|----------|-----------------|----------|
| WO 2003050295   | A2   | 20030619 | WO 2002-US39993 | 20021212 |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM         | RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG     |          |                 |          |
| WO 2003066005   | A2   | 20030814 | WO 2003-US4283  | 20030210 |
| WO 2003066005   | A3   | 20040610 |                 |          |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM     | RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG |          |                 |          |
| WO 2004054624   | A1   | 20040701 | WO 2003-US18776 | 20030612 |
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NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,  
GW, ML, MR, NE, SN, TD, TG

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| PRIORITY APPLN. INFO.: | US 2001-340762P | P 20011212 |
|                        | US 2002-355275P | P 20020208 |
|                        | US 2002-367055P | P 20020322 |
|                        | WO 2002-US39993 | A 20021212 |

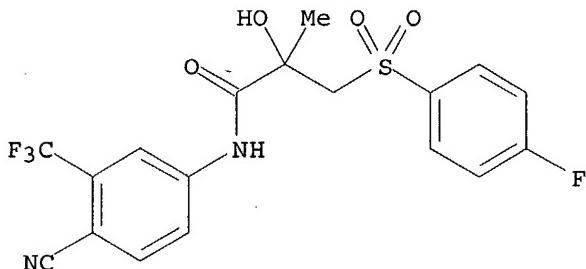
OTHER SOURCE(S): CASREACT 139:30801

IT 90357-06-5, Bicalutamide

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL  
(Biological study); USES (Uses)  
(assays and implements for determining and modulating heat shock protein 90  
binding activity, and therapeutic use with other agents)

RN 90357-06-5 CAPLUS

CN Propanamide, N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-fluorophenyl)sulfonyl]-2-hydroxy-2-methyl- (9CI) (CA INDEX NAME)



AB Ligand binding assays as applied to HSP90s as receptors or ligands, and reagents useful therefore, are described and claimed, as are methods of assaying for HSP90 modulators and methods of using the resulting products identified thereby. The methodol. of the invention may be used in the treatment and prevention of an HSP90-mediated disease, e.g. cancer. Modulators of the invention include e.g. ansamycins.

L7 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 2

ACCESSION NUMBER: 2002:964133 CAPLUS

DOCUMENT NUMBER: 138:24551

TITLE: Preparation of rac-bicalutamide

INVENTOR(S): Dolitzky, Ben-Zion; Reany, Ofer; Shamai, Jenny

PATENT ASSIGNEE(S): Teva Pharmaceutical Industries Ltd., Israel; Teva Pharmaceuticals USA, Inc.; Biogal Gyogyszergyar

SOURCE: PCT Int. Appl., 22 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

| PATENT NO.  | KIND | DATE     | APPLICATION NO. | DATE     |
|---|------|----------|-----------------|----------|
| WO 2002100339   | A2   | 20021219 | WO 2002-US18329 | 20020613 |
| WO 2002100339   | A3   | 20031016 |                 |          |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM |      |          |                 |          |
| RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,   |      |          |                 |          |

Priority  
JULY 10 2002

CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,  
 BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG  
 EP 1406855 A2 20040414 EP 2002-739801 20020613  
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 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR  
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 US 2002-371069P P 20020409 ✓  
 WO 2002-US18329 W 20020613

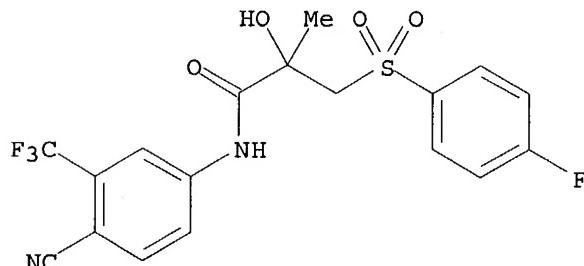
OTHER SOURCE(S): CASREACT 138:24551

IT 90357-06-5P, Bicalutamide

RL: SPN (Synthetic preparation); PREP (Preparation)  
 (preparation of rac-bicalutamide)

RN 90357-06-5 CAPLUS

CN Propanamide, N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-fluorophenyl)sulfonyl]-2-hydroxy-2-methyl- (9CI) (CA INDEX NAME)



AB Racemic and optically active N-[4-cyano-3-trifluoromethylphenyl]-3-[4-fluorophenylsulfonyl]-2-hydroxy-2-Me propionamide (bicalutamide) were prepared starting from Et pyruvate and Me methacrylate. Thus, 5-amino-2-cyanobenzotrifluoride was treated with DABCO and reacted with deprotonated ethyl-[2-(4-fluorophenyl sulfone)]-2-hydroxy propionate (prepared from Et pyruvate) to give ~40% rac-bicalutamide. **Micronized** particles of rac-bicalutamide can be obtained as pharmaceutical compns. that are useful for its anti-androgen activity (no data). Bicalutamide intermediates were also prepared, including ethyl-[2-(4-fluorophenyl sulfone)]-2-hydroxy propionate, 1,2-epoxy-2-Me propionate and 2-hydroxy-2-methyl-3-(4-fluorophenylthio) propionic acid.

L7 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2002:637506 CAPLUS

DOCUMENT NUMBER: 137:190729

TITLE: Novel modified-release formulation containing amphiphilic lipids as a hydrophobic matrix former

INVENTOR(S): Juppo, Anne

PATENT ASSIGNEE(S): Astrazeneca Ab, Swed.

SOURCE: PCT Int. Appl., 31 pp.  
 CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

| PATENT NO.   | KIND | DATE     | APPLICATION NO. | DATE     |
|--|------|----------|-----------------|----------|
| WO 2002064121  | A1   | 20020822 | WO 2002-SE228   | 20020208 |
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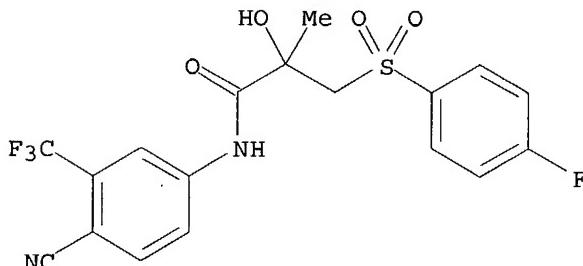
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 EP 1368006 A1 20031210 EP 2002-710645 20020208  
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IE, SI, LT, LV, FI, RO, MK, CY, AL, TR  
 BR 2002006825 A 20040225 BR 2002-6825 20020208  
 JP 2004518709 T2 20040624 JP 2002-563916 20020208  
 US 2004067256 A1 20040408 US 2003-467900 20030811  
 NO 2003003564 A 20031002 NO 2003-3564 20030812  
 PRIORITY APPLN. INFO.: SE 2001-477 A 20010213  
 SE 2001-478 A 20010213  
 WO 2002-SE228 W 20020208

IT 90357-06-5, Bicalutamide

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
(controlled-release multiparticulate solid dispersion formulation  
containing amphiphilic lipid matrix)

RN 90357-06-5 CAPLUS

CN Propanamide, N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-  
fluorophenyl)sulfonyl]-2-hydroxy-2-methyl- (9CI) (CA INDEX NAME)



AB The present invention is directed to a multiparticulate, modified release solid dispersion formulation, comprising a drug substance having a water-solubility of  $\leq$  8 mg/mL at room temperature, a hydrophobic matrix former which is a water insol., non-swelling amphiphilic lipid, and a hydrophilic matrix former which is a meltable, water-soluble excipient. The weight ratio of

hydrophobic matrix former to hydrophilic matrix former is  $\geq$  1, and the particle size is  $<$  300  $\mu\text{m}$ . Also a unit dosage of the same, as well as process for the preparation thereof and the use of the formulation and unit dosage are claimed. For example, felodipine (1 g) was dissolved in a melt of 4 g cetanol at 110° and 2 g of Poloxamer 407 was added into the melt. The melted mixture was kept at 110° and atomized at air temperature of 400° and a pressure of 7 bar; the particles were collected into a vessel which was kept at temperature -50°, and thereafter dried over night in a vacuum oven at 25° and 2 mbar. The resulted particles had a 90% fractile size (90% smaller than) of 77  $\mu\text{m}$  and a roundness of 0.87.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2002:293419 CAPLUS  
 DOCUMENT NUMBER: 136:315004  
 TITLE: Liposomes encapsulating anticancer drugs and the use thereof in the treatment of malignant tumors  
 INVENTOR(S): Parente Duena, Antonio; Pons Lambiez, Ferran; Fabra Fres, Angels; Polo Trasancos, Maria Dolores; Garcés

PATENT ASSIGNEE(S) : Garces, Josep; Reig Isart, Francesca  
 SOURCE: Lipotec, S.A., Spain  
 PCT Int. Appl., 38 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Spanish  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

| PATENT NO.   | KIND | DATE     | APPLICATION NO. | DATE       |
|--|------|----------|-----------------|------------|
| WO 2002030397  | A1   | 20020418 | WO 2001-ES367   | 20011003   |
| WO 2002030397  | C2   | 20040513 |                 |            |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,<br>CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,<br>GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,<br>LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL,<br>PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,<br>US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM<br>RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,<br>DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,<br>BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG |      |          |                 |            |
| ES 2186484   | A1   | 20030501 | ES 2000-2447    | 20001010   |
| ES 2186484   | B1   | 20040701 |                 |            |
| AU 2001091904  | A5   | 20020422 | AU 2001-91904   | 20011003   |
| EP 1325739   | A1   | 20030709 | EP 2001-972111  | 20011003   |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,<br>IE, SI, LT, LV, FI, RO, MK, CY, AL, TR   |      |          |                 |            |
| BR 2001013552  | A    | 20030729 | BR 2001-13552   | 20011003   |
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| PRIORITY APPLN. INFO.:   |      |          | ES 2000-2447    | A 20001010 |
|  |      |          | WO 2001-ES367   | W 20011003 |

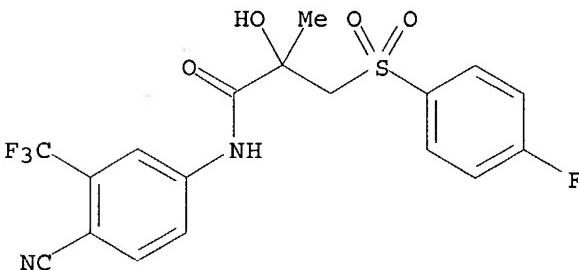
IT 90357-06-5, Bicalutamide

RL: PEP (Physical, engineering or chemical process); PYP (Physical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)

(lipopeptide liposomes encapsulating anticancer drugs and the use thereof in the treatment of malignant tumors)

RN 90357-06-5 CAPLUS

CN Propanamide, N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-fluorophenyl)sulfonyl]-2-hydroxy-2-methyl- (9CI) (CA INDEX NAME)



AB The invention relates to liposomes encapsulating anticancer drugs and the use thereof in the treatment of malignant tumors. The liposomes are covered by a lipopeptide consisting of three substructures: a lipid fragment, an active oligopeptide and a spacer oligopeptide between the two fragments. Said liposomes can be used in i.v. administration for the treatment of malignant tumors.

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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|--|------------|---------|
| => log Y                                   |            |         |
| COST IN U.S. DOLLARS                       | SINCE FILE | TOTAL   |
|  | ENTRY      | SESSION |
| FULL ESTIMATED COST                        | 34.16      | 39.64   |
| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE | TOTAL   |
|  | ENTRY      | SESSION |
| CA SUBSCRIBER PRICE                        | -3.50      | -3.50   |

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